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Reply

Dr. Laoutaris and colleagues believe that high-intensity endurance inspiratory muscle training (IMT) with the TRAINAIR device (Project Electronics Ltd., London, United Kingdom), as used in their nonrandomized studies (1–3), is associated with improvement in peak oxygen consumption (VO_2), while IMT using the threshold device (Threshold Inspiratory Muscle Trainer, Healthscan Products Inc., Cedar Grove, New Jersey) at 30% of maximal inspiratory pressure (PI_{max}), as used in our controlled experiments (4,5), may not increase peak VO_2 in patients with chronic heart failure. They failed to note that we also measured PI_{max} during an incremental test and endurance time (4), measurements which they believe to be associated with changes in peak VO_2 . In agreement with our findings, other investigators (6) have shown early improvement in PI_{max} and exercise capacity after IMT with the threshold device, and the data of Laoutaris et al. (1) show a 28% increment in PI_{max} after only 4 weeks of IMT (Fig. 3 of Laoutaris et al. [1]). Moreover, in our mechanistic study on patients with heart failure (5), improvement in PI_{max} was correlated with diaphragm hypertrophy, in contrast to their beliefs based on experiments using rat skeletal muscle or inspiratory muscles of normal individuals.

The improvement in ventilatory response to exercise in our study comes as no surprise, since we have recently shown that inspiratory muscle weakness is associated with augmented peripheral chemoreflex response, a major determinant of ventilatory efficiency (7). Therefore, the finding of Laoutaris et al. that ventilatory efficiency did not improve with IMT is probably explained by the inclusion of 79% of patients without inspiratory muscle weakness (3), a bias that may explain the negative results of other trials.

Contrary to the beliefs of Dr. Laoutaris and colleagues, it is the evidence in support of the efficacy of their high intensity endurance IMT in patients with heart failure without inspiratory muscle weakness that is frail. In their efficacy study (1), no correction for multiple comparisons was used, but when the appropriate statis-

tical analysis was applied in their mechanistic studies, no interaction was found for the changes in PI_{max} or in peak VO_2 (2,3). Finally, despite the fact that our controlled (4,5) experiments and the data of Laoutaris et al. (1–3) disprove their beliefs, we consider that they have raised interesting hypotheses that must be tested by well-designed research.

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